IN THE CLAIMS:

This listing of the claims will replace all prior versions and listing of the claims in the present application:

- 1. (Currently Amended.) A method of treating an intestinal damage comprising administering a pharmaceutically active formulation of PYY or a PYY agonist to a subject <u>human</u> to treat the intestinal damage.
- 2. (Previously Presented.) The method of claim 1 wherein the intestinal damage is associated with a condition selected from the group consisting of inflammatory bowel disease, bowel atrophy, loss of bowel mucosa, and loss of bowel mucosal function.
- 3. (Previously Presented.) The method of claim 2 wherein the inflammatory bowel disease is ulcerative colitis.
- 4. (Cancelled.)
- 5. (Previously Presented.) The method of claim 1 wherein the intestinal damage is caused by an event selected from the group consisting of exposure to cytotoxic agents, radiation, toxicity, infection and an injury.
- 6. (Previously Presented.) The method of claim 1, wherein the PYY or the PYY agonist is used in conjunction with a cytotoxic agent or radiation.
- 7. (Withdrawn.) The method of claim 1 further comprising administering a growth hormone.
- 8. (Previously Presented.) The method of claim 1 further comprising administering an antiinflammatory agent.
- 9. (Previously Presented.) The method of claim 8 wherein the anti-inflammatory agent is selected from the group consisting of tacrolimus, mycophenolate mofetil, anti-tumor necrosis factor antibody, interleukin-10, interleukin-11, anti-interleukin-12 antibody, anti-inlerleukin-1 antibody, anti-alpha4 integrin antibody, and nicotine.

- 10. (Previously Presented.) The method of claim 1 wherein the PYY or the PYY agonist is administered by a route selected from the group consisting of intravenous, intraperitoneal, subcutaneous, intramuscular, oral, rectal, topical, transmucosal, nasal, or pulmonary inhalation.
- 11. (Previously Presented.) The method of claim 1 wherein the PYY or the PYY agonist is administered in the amount of about 100 µg to about 500 mg/day.
- 12. (Previously Presented.) The method of claim 1 wherein the PYY or the PYY agonist is administered in the amount of about 500 µg to 100 mg/day.
- 13. (Cancelled.)
- 14. (Previously Presented.) The method of claim 1 wherein the PYY agonist is PYY[3-36].
- 15. (Withdrawn.) A probiotic bacterium comprising a nucleic acid encoding PYY or a PYY agonist.
- 16. (Withdrawn.) The probiotic bacterium of claim 15 wherein the bacteria expresses and secretes the PYY or the PYY agonist.
- 17. (Withdrawn.) The probiotic bacterium of claim 15 wherein the bacterium is a lactobacillus bacterium.
- 18. (Withdrawn.) The probiotic bacterium of claim 15 wherein the PYY agonist is PYY[3-36].
- 19. (Withdrawn.) A method of treating a bowel condition comprising administering the probiotic bacterium of claim 15 to a patient.
- 20. (Withdrawn.) The method of claim 19 wherein the probiotic bacterium is administered by oral ingestion or suppository.
- 21. (Withdrawn.) The method of claim 19 wherein the bowel condition comprises intestinal damage.